The listing of claims will replace all prior versions, and listings, of claims in the application: <u>Listing of Claims</u>:

1. (Previously Presented) A compound of formula (I) or a salt, N-oxide, hydrate or solvate thereof:

$$\begin{array}{c} R_1 \\ R_2 \\ HN \\ N \end{array} \begin{array}{c} R_2 \\ R_3 \end{array}$$
 (I)

wherein

R₁ is a group of formula (IA):

$$-Ar^{1}$$
- $(Alk^{1})_{p}$ - $(Z)_{r}$ - $(Alk^{2})_{s}$ - Q (lA)

wherein in any compatible combination

Ar¹ is an optionally substituted aryl or heteroaryl radical,

 Alk^{1} and Alk^{2} are optionally substituted divalent C_{1} - C_{6} alkylene or C_{2} - C_{6} alkenylene radicals.

p, r and s are independently 0 or 1,

- R_2 is (i) a group of formula (IA) as defined in relation to R_1 ;
 - (ii) a carboxamide radical; or
 - (iii) a non aromatic carbocyclic or heterocyclic ring wherein a ring carbon is optionally substituted, and/or a ring nitrogen is optionally substituted by a group of formula -

(Alk¹)_p-(Z)_r-(Alk²)-Q wherein Q, Alk¹, Alk², Z, p, r and s are as defined above in relation to group (IA); and

R₃ is hydrogen, or methyl, ethyl, n- or iso-propyl any of which being optionally substituted by hydroxy;

X is $-OR_4$ or $-NR_4R_5$ wherein R_4 and R_5 independently represent hydrogen or optionally substituted C_1 - C_6 alkyl, or R_4 and R_5 taken together with the nitrogen to which they are attached form an optionally substituted nitrogen-containing ring having 5-8 ring atoms.

2. (Previously Presented) The compound as claimed in claim 1 wherein in the compound of formula (I), R_1 has formula (IB):

$$R_{e}$$
 OH
 (IB)

wherein R_6 is chloro, bromo, C_1 - C_6 alkyl, or cyano.

3. (Previously Presented) The compound as claimed in claim 1 wherein in the compound of formula (I) R_1 has formula (IC):

$$Q-(Alk^2)_s-(Z)_r-(Alk^1)_p$$

$$OH$$
(IC)

wherein Alk¹, Alk², p, r, s, Z and Q are as defined in claim 1 in relation to formula (IA), and R

represents one or more optional substituents.

- 4. (Withdrawn) The compound as claimed in claim 2 wherein R is -OH in the 4- position of the phenyl ring and the $-(Alk^1)_p-(Z)_r-(Alk^2)_s$ -Q substituent is in the 5- position of the phenyl ring.
- 5. (Withdrawn) The compound as claimed in claim 4 wherein r is 0, and Q is hydrogen or optionally substituted phenyl.
- 6. (Withdrawn) The compound as claimed in claim 5 wherein s is 0, p is 1 and Alk¹ is a nonsubstituted divalent C_1 - C_6 alkylene or C_2 - C_6 alkenylene radical.
- 7. (Withdrawn) The compound as claimed in claim 5 wherein Alk¹ is -CH₂-, -CH₂CH₂-, -CH₂CH₂-, or -CH=CH-.
- 8. (Withdrawn) The compound as claimed in claim 4 wherein p, r and s are each 0.
- 9. (Previously Presented) The compound as claimed in claim 1 wherein R_2 is phenyl, 2-, 3-, or 4-pyridyl, 2- or 3-furanyl, 2- or 3-thienyl, or thiazolyl, optionally substituted by one or more of methoxy, ethoxy, methylenedioxy, ethylenedioxy, fluoro, chloro, bromo, or trifJuoromethyl.
- 10. (Previously Presented) The compound as claimed in claim 1 wherein R_2 is optionally substituted phenyl.
- (Previously Presented) The compound as claimed in claim 1 wherein R₂ is phenyl substituted in the 4 position by (i) C₁-C₆ alkoxy such as methoxy or ethoxy, fluoro, chloro, bromo, morpholinomethyl, piperazino, N-methylpiperazino, or piperidino, (ii)optionally substituted C₁-C₆ alkyl, eg optionally substituted methyl, ethyl, n-propyl or iso-propyl (iii) optionally substituted morpholino C₁-C₆ alkyl-, thiomorpholino C₁-C₆ alkyl-, piperazino C₁-C₆ alkyl-, methyl piperazino C₁-C₆ alkyl-, or diethylamino (iv) -NH₂, -NHR^A, -NR^AR^B, -NHCOR^A, -NHCOR^A, -NHCOR^A, -NHCOR^A, -NHSO₂0R^A, -NR^BSO₂OR^A,

-NHCONH₂, -NR^ACONH₂, NHCONHR^B, -NR^ACONHR^B, -NHCONR^AR^B, or -NR^ACONR^AR^B wherein R^A and R^B are independently a (C_1-C_6) alkyl group or (v) optionally substituted piperadino, piperazino, morpholino or thiomorpholino.

12. (Withdrawn) The compound as claimed in claim 1 wherein R_2 is a carboxamide radical of formula $-CONR^B(Alk)_nR^A$ wherein

Alk is an optionally substituted divalent alkylene, alkenylene or alkynylene radical,

n is 0 or 1,

R^B is hydrogen or a C₁-C₆ alkyl or C₂-C₆ alkenyl group,

R^A is hydroxy or an optionally substituted carbocyclic or heterocyclic ring,

or R^A and R^B taken together with the nitrogen to which they are attached form an N-heterocyclic ring which may optionally contain one or more additional hetero atoms selected from O, S and N, and which may optionally be substituted on one or more ring C or N atoms.

13. (Withdrawn) The compound as claimed claim 12 wherein

Alk is an optionally substituted –CH₂-, -CH₂CH₂-, -CH₂CH₂-, -CH₂CH₂-, -CH₂CH=CH-, or –CH₂CCCH₂- radical.

n is 0 or 1,

R^B is hydrogen, methyl, ethyl, n- or iso-propyl, or allyl,

R^A is hydroxy, hydroxy and/or chloro-substituted phenyl, 3,4 methylenedioxyphenyl,

pyridyl, furyl, thienyl, N-piperazinyl, or N-morpholinyl,

or R^A and R^B taken together with the nitrogen to which they are attached form a morpholino, piperidinyl, piperazinyl or N-phenylpiperazinyl ring.

- 14. (Withdrawn) The compound as claimed in claim 12 wherein n is 0, R^B is hydrogen and R^A is hydroxy or an optionally substituted carbocyclic or heterocyclic ring.
- 15. (Withdrawn) The compound as claimed in claim 1 wherein R₃ is hydrogen.
- 16. (Previously Presented) The compound as claimed in claim 1 wherein R₃ is other than hydrogen and the stereochemical configuration at the carbon centre to which it is attached is that of a D amino acid.
- 17. (Previously Presented) The compound as claimed in claim 1 wherein X is $-OR_4$ or $-NHR_4$ wherein R_4 is C_1 - C_6 alkyl, optionally substituted by hydroxy, or a primary- secondary, tertiary- or cyclic-amino group
- 18. (Withdrawn) The compound as claimed in claim 1 wherein X is $-NR_4R_5$ wherein R_4 and R_5 taken together with the nitrogen to which they are attached form a morpholino, piperidinyl or piperazinyl ring, the latter being optionally substituted by C_1 - C_6 alkyl on the second nitrogen.
- 19. (Withdrawn) A method of treatment of diseases or conditions mediated by excessive or inappropriate HSP90 activity in mammals which method comprises administering to the mammal an amount of a compound of formula (I) as defined in claim 1, or a salt, hydrate or solvate thereof, effective to inhibit said HSP90 activity.
- 20. (Withdrawn) The method as claimed claim 19 for immunosupression or the treatment of cancer; viral disease, inflammatory diseases such as rheumatoid arthritis, asthma, multiple sclerosis, Type I diabetes, lupus, psoriasis and inflammatory bowel disease; cystic fibrosis

angiogenesis-related disease such as diabetic retinopathy, haemangiomas, and endometriosis; or for protection of normal cells against chemotherapy-induced toxicity; or diseases where failure to undergo apoptosis is an underlying factor; or protection from hypoxia-ischemic injury due to elevation of Hsp70 in the heart and brain; scrapie/CJD, Huntingdon's and Alzheimer's disease.

- 21. (Canceled)
- 22. (Canceled)
- 23. (Previously Presented) A compound

or a salt, solvate or hydrate thereof.

24. (Previously Presented) A pharmaceutical or veterinary composition comprising a compound as defined in claim 1, together with a pharmaceutically or veterinarily acceptable carrier.